

Exagon 400 mg/ml solution for injection

Atļautas

- Pentobarbital sodium

Zāļu identifikācija

Zāļu nosaukums:

Exagon 400 mg/ml solution for injection

Aktīvā viela:

Pieejamas tikai [English](#)

Mērķsugas:

Suns

Kaķis

Zaķis

Trusis

Jūdescūciņa

Kāmis

Pele

Žurka

Balodis

Liellops

Mājputni

Zirgs

Zirgs (ponijs)

Ūdele

Sesks

Cūka
Putni
Varde
Ķirzaka
Čūska
Ūdens bruņurupucis

Lietošanas veids:

intraperitoneālai lietošanai
intravenozai lietošanai
intrakardiālai lietošanai
Intrapulmonālai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai [English](#)
400.00 miligrams(i) / 1.00 mililitrs(i)

Farmaceutiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

intraperitoneālai lietošanai:

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Zaķis

- Gaļa un blakusprodukti. no withdrawal period

Do not use in animals intended for human or animal consumption. Adequate measures must be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

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Trusis

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intravenozai lietošanai:

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Balodis

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- Ola. no withdrawal period

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Liellops

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- Piens. no withdrawal period

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Mājputni

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Zirgs

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Zirgs (ponijs)

- Piens. no withdrawal period

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Cūka

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intrakardiālai lietošanai:

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Zaķis

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Cūka

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Intrapulmonālai lietošanai:

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Balodis

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- Ola. no withdrawal period

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used for human or animal consumption.

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Mājputni

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Do not use in animals intended for human or animal consumption. Adequate measures must be taken to ensure that carcasses of animals treated with this product and the by-products of these animals do not enter the food chain and are not used for human or animal consumption.

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Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QN51AA01

Izplatīšanas juridiskie nosacījumi:

Recepšu veterinārās zāles

Atļaujas statuss:

Derīga

Atļautas:

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Pieejams:

Finland

Iepakojuma apraksts:

Pieejamas tikai [English](#)

Pieejamas tikai [English](#)

Papildu informācija

Tiesību tips:

Marketing Authorisation

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:

Pieteikums ģenēriskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(1) pants)

Tirdzniecības atļaujas turētājs:

Vetviva Richter GmbH

Tirdzniecības atļaujas datums:

9/04/2014

Ražošanas vietas sēriju izlaidei:

Vetviva Richter GmbH

Atbildīgā iestāde:

Finnish Medicines Agency

Atļaujas numurs:

31523

Atļaujas statusa maiņas datums:

9/04/2014

Atsauces dalībvalsts:

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Procedūras numurs:

DE/V/0155/001

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Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz:
www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Zāļu apraksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Lietošanas instrukcija

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

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